## Amendments to the Claims

The Listing of Claims presented below will replace all prior versions, and listings, of claims in the application.

## Listing of Claims

A mixture or set of sub-mixtures comprising X-mer precursors,
wherein the X-mer precursors have a minimum length of 3 nucleotides;
wherein the mixture has a minimum mixture coverage complexity of at least
56/N or wherein the set of sub-mixtures has a composite mixture coverage

complexity of at least 56/N, wherein N represents the number of distinct X-mer

precursors in the mixture;

wherein each sub-mixture in said set has a reduced mixture coverage complexity as compared with the composite mixture coverage complexity;

wherein each sub-mixture comprises a plurality of X-mer precursors; wherein said length is selected independently for each X-mer precursor; and wherein the mixture or set of sub-mixtures further comprises a set of tags wherein each tag is covalently linked to at least one X-mer precursor through a cleavable linker.

2. A mixture or set of sub-mixtures comprising X-mer precursors,

wherein said X-mer precursors have a minimum length of 3 nucleotides; wherein said mixture has a minimum mixture coverage complexity of at least 56/N or wherein said set of sub-mixtures has a composite mixture coverage complexity of at least 56/N, wherein N represents the number of distinct X-mer precursors in the mixture;

wherein each sub-mixture in said set has a reduced mixture coverage complexity as compared with the composite mixture coverage complexity;

wherein each sub-mixture further comprises a plurality of X-mer precursors; wherein said length is selected independently for each X-mer precursor;

Atty Ref No.: 2003309(AGIL)-0013 Client Ref. No.: 10992153-1 wherein the mixture or set of sub-mixtures further comprises a set of tags wherein each tag is covalently linked to at least one X-mer precursor through a cleavable linker; and

wherein said X-mer precursors have a determined isotopic composition.

- 3. The mixture or set of sub-mixtures of claim 1 or 2 wherein said mixture has a mixture coverage complexity of at least about 1/2 when said mixture contains at least 128 discrete X-mers, or wherein said set of sub-mixtures has a composite mixture coverage complexity of at least about 1/2 when said set of sub-mixtures contains at least 128 discrete X-mers.
- 4. The mixture or set of sub-mixtures of claim 1 or 2, wherein said mixture has a mixture coverage complexity of at least about 1/4 when said mixture contains at least 256 discrete X-mers, or wherein said set of sub-mixtures has a composite mixture coverage complexity of at least about 1/4 when said set of sub-mixtures contains at least 256 discrete X-mers.
- 5. The mixture or set of sub-mixtures of claim 1 or 2, wherein said mixture has a mixture coverage complexity of at least about 1/8 when said mixture contains at least 512 discrete X-mers, or wherein said set of sub-mixtures has a composite mixture coverage complexity of at least about 1/8 when said set of sub-mixtures contains at least 512 discrete X-mers.
- 6. The mixture or set of sub-mixtures of claim 1 or 2, wherein nucleotide sequences of the precursors of said mixture or set of sub-mixtures are known.
- 7. The mixture or set of sub-mixtures of claim 1 or 2, wherein a number of tags in the set of tags distinguishable by mass spectrometry after cleavage of the linkers is between approximately 10-100,000.

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- 8. The mixture or set of sub-mixtures of claim 1 or 2, wherein a number of tags in the set of tags distinguishable by mass spectrometry after cleavage of the linkers is between approximately 20-20,000.
- 9. The mixture or set of sub-mixtures of claim 1 or 2, wherein a number of tags in the set of tags distinguishable by mass spectrometry after cleavage of the linkers is between approximately 20-10,000.
- 10. The mixture or set of sub-mixtures of claim 1 or 2, wherein a number of tags in the set of tags distinguishable by mass spectrometry after cleavage of the linkers is between approximately 20-5,000.
- 11. The mixture or set of sub-mixtures of claim 1 or 2, wherein a number of tags in the set of tags distinguishable by mass spectrometry after cleavage of the linkers is between approximately 50-1000.
- 12. The mixture or set of sub-mixtures of claim 1 or 2, wherein a number of tags in the set of tags distinguishable by mass spectrometry after cleavage of the linkers is greater than a mass number complexity (MNC) of a natural equivalent of the mixture or set of sub-mixtures, wherein the natural equivalent of the X-mer precursors are extended by one nucleotide, and wherein the number of tags in the set of tags is less than or equal to a number of X-mer precursors in the mixture or set of sub-mixtures.
- 13. The mixture or set of sub-mixtures of claim 1 or 2, wherein a number of tags in the set of tags distinguishable by mass spectrometry after cleavage of the linkers is greater than 75% of a mass number complexity (MNC) of a natural equivalent of mixture or set of sub-mixtures, wherein the natural equivalent of the X-mer precursors are extended by one nucleotide, and wherein the number of tags in the set of tags is less than or equal to a number of X-mer precursors in the mixture or set of sub-mixtures.

14. The mixture or set of sub-mixtures of claim 1 or 2, wherein a number of tags in the set of tags distinguishable by mass spectrometry after cleavage of the linkers is at least 0.5% of a number of X-mer precursors in the mixture or set of sub-mixtures,

and less than or equal to the number of X-mer precursors in the mixture or set of sub-

mixtures.

15. The mixture or set of sub-mixtures of claim 1 or 2, wherein a number of tags in the

set of tags distinguishable by mass spectrometry after cleavage of the linkers is at

least 1% of a number of X-mer precursors in the mixture or set of sub-mixtures, and

less than or equal to the number of X-mer precursors in the mixture or set of sub-

mixtures.

16. The mixture or set of sub-mixtures of claim 1 or 2, wherein a number of tags in the

set of tags distinguishable by mass spectrometry after cleavage of the linkers is at

least 10% of a number of X-mer precursors in the mixture or set of sub-mixtures, and

less than or equal to the number of X-mer precursors in the mixture or set of sub-

mixtures.

17. The mixture or set of sub-mixtures of claim 1 or 2, wherein a number of tags in the

set of tags distinguishable by mass spectrometry after cleavage of the linkers is at

least 25% of a number of X-mer precursors in the mixture or set of sub-mixtures, and

less than or equal to the number of X-mer precursors in the mixture or set of sub-

mixtures.

18-73. (**Cancelled**)

- 74. A kit for carrying out a method of analyzing a target nucleic acid sequence, comprising:
  - a. the mixture or the set of sub-mixtures of claim 1; and
  - b. an enzyme having a nucleotide polymerase activity.

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- 75. The kit of claim 74, further comprising a multiplicity of nucleotides selected from the group consisting of natural chain-terminating triphosphates and modified chain-terminating triphosphates.
- 76. The kit of claim 74, further comprising chain-terminating nucleotides with an affinity label for purification of nucleic acids.
- 77. A kit for carrying out a method of analyzing a target nucleic acid sequence comprising:
  - a. the mixture or the set of sub-mixtures of claim 1; and
  - b. a DNA ligase.
- 78. A kit for carrying a method of analyzing a target nucleic acid sequence, comprising:
  - a. the mixture or the set of sub-mixtures of claim 1; and
  - b. a condensing agent.
- 79. A kit for carrying out a method of analyzing a target nucleic acid sequence having a 3'-end and a 5'-end, comprising:
  - a. the mixture or the set of sub-mixtures of claim 1;
  - b. a DNA ligase; and
  - c. an array comprising:
    - (a) a surface; and
    - (b) a multiplicity of nucleic acid sequence probes comprising:
      - (i) a nucleic acid attached to said surface, wherein the nucleic acid has a terminal 3'-hydroxyl end and wherein the 5' end is directly or indirectly attached to said surface.
- 80. A kit for carrying out a method of analyzing a target nucleic acid sequence having a 3'-end and a 5'-end, comprising:
  - a. the mixture or the set of sub-mixtures of claim 1;
  - b. a condensing agent; and
  - c. an array comprising:

- (a) a surface; and
- (b) a multiplicity of nucleic acid sequence probes comprising:
- (i) a nucleic acid attached to said surface, wherein the nucleic acid has a terminal 3'-hydroxyl end and wherein the 5' end is directly or indirectly attached to said surface.